5

10

20

35

What is claimed is:

## CLAIMS



- Monoclonal antibodies or their fragments, more particularly their Fv, Fab, and F(ab')2 fragments, characterized in that they recognize an epitope of a bacterium of the species T. equigenitalis.
- 2/ Monoclonal antibodies or their fragments, more particularly their Fv, Fab, and F(ab')2 fragments, according to claim 1 characterized in that they do not exhibit a crossed reaction with the epitope or epitopes of a bacterium of a different Taylore la species or of a bacterium of a different genus.
- 3/ Monoclonal antibodies or their fragments, according to claim 1 or 2, characterized in that they are capable of 15 recognizing proteins of *T. equigenitalis* of the group comprising proteins such as proteins of 150 kDa, 120 kDa, 52.7 kDa or 22 (LPS) kDa.
  - 4/ Monoclonal antibodies, characterized in that they can be obtained from hybrids
  - by fusion of non-secreting murine myeloma cells with spleen cells from mice immunized using an inactivated strain of the species T. equipmentalis of extract(s) from such a strain, and
- cloning and selection according to the capacity of their culture supernatant to recognize an epitope or epitopes of a bacterium of the species T. equigenitalis,
  - recovery of the required monoclonal antibodies, followed by purification if necessary.
- 5/ Immunogenic proteins, characterized in that they are 30 capable of interacting with monoclonal antibodies or their fragments according to any one of claims 1 to 4.
  - 6/ Monoclonal antibodies, and their fragments, in particular their Fv, Fab, F(ab') fragments, characterized in that they are anti-antibodies, i.e. antibodies capable of interacting with the monoclonal antibodies or their fragments according to any one of claims 1 to 4.
  - 7/ A method of obtaining monoclonal antibodies according to any one of claims 1 to 4, characterized in that it

## comprises:

5

10

15

20

30

35

- fusion of non-secreting murine myeloma cells with spleen cells from mice immunized by means of a strain of the species *T. equigenitalis* or extract(s) from such a strain,
- screening by means of a detection technique, such as in particular indirect immunofluorescence, of hybridomas whose culture supernatants exhibit a positive reaction with a bacterium of the species *T. equigenitalis* or a fragment of the latter,
- selection by cloning of these hybridomas with respect to their reactivity, in relation to *T. equigenitalis*, and
  - recovery of the monoclonal antibodies, followed if necessary by their purification.
  - 8/ A method of obtaining monoclonal antibodies according to claim 6, characterized in that it comprises:
  - fusion of non-secreting murine myeloma cells with spleen cells from mice immunized using monoclonal antibodies or their fragments as defined in one of claims 1 to 4,
  - screening by means of a detection technique, such as in particular indirect immunofluorescence, of hybridomas whose culture supernatants exhibit a positive reaction with one of the said monoclonal antibodies or their fragments,
    - selection by cloping of these hybridomas, and
    - recovery of the required anti-antibodies.
- 9/ Strains of hybridomas, characterized in that they are capable of secreting monoclonal antibodies according to any one of claims 1 to 4.
  - 10/ Strains of hybridomas, characterized in that they are capable of secreting monoclonal antibodies according to claim 6.
  - 11/ Method of identification of a bacterium of the species *T. equigenitalis* in a sample or in a culture, comprising:
  - bringing the sample or the culture to be analysed, which may contain *T. equigenitalis*, into contact with
  - i. an effective quantity of at least one monoclonal antibody or a fragment of such an antibody according to any one of claims 1 to 4 and, optionally, blocking the non

5

15

20

antigen-antibody reactions,

ii. or, as a variant, to detect the presence of antibodies directed against T. equipmentalis with an immunogenic protein according to claim 5 or an antibody according to claim 6,

in conditions permitting \( \alpha \) reaction of the antigen-antibody type and

- detection of any product formed in a reaction of the antigen-antibody type.
- 10 12/ Method of diagnosis of an infection by T.

  equigenitalis, more particularly contagious equine metritis
  in a sample or a culture, comprising:
  - bringing one or more monoclonal antibodies according to any one of claims 1 to 4 or their fragments, into contact with a biological sample, and
  - detection of the reaction of the antigen-antibody type produced in the case when *T* equigenitalis is present in the sample,
  - and, optionally, blocking of the non antigen-antibody reactions, for example, by saturation of the specimen obtained by means of a serum from which anti-T. equigenitalis antibodies have been removed.
  - 13/ Kits for the application of a method according to one of claims 11 or 12, characterized in that they include
- one or more monoclonal antibodies, or their fragments, according to any one of claims 1 to 4, or at least one immunogenic protein according to claim 5, or one or more monoclonal antibodies, or their fragments, according to claim 6,
- reagents, in particular markers or buffers, for carrying out the intended immunogenic reaction, and, optionally, reagents for blocking non antigen-antibody reactions such as mouse serum,
  - as well as instructions for use.
- 14/ Pharmaceutical compositions, characterized in that they contain one or more monoclonal antibodies, or their fragments, according to any one of claims 1 to 4, as vectors of medicaments or as agents for passive immunotherapy, alone

5

or in combination with pharmaceutically inert vehicles.

15/ Vaccinal compositions, characterized in that they combination with physiologically acceptable contain, least immunogenic protein as defined excipients, at \_ong  $\sqrt{\text{or/one}}$  antibody according to claim 6, according to claim 5, or one fragment of one such antibody, in sufficient quantity to evoke an immune response.

16/ Use of the monoclonal antibodies according to one of

4 for the elaboration of biosensors.